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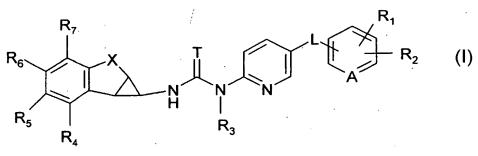
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(54) Title: NON-NUCLEOTIDE REVERSE TRANSCRIPTASE INHIBITORS



(57) Abstract: Compounds of the formula Z: where, A is CH or N; R₁ is a substituent to a carbon atom in the ring containing A selected from -S(=O)_pRa, where Ra is -C₁-C₄ alkyl, -ORx, -NRxRx, -NHNRxRx, - NHNHC(=O)ORx, -NRxOH; -C(=O)-Rb, where Rb is -CT-C4-alkyl, ORx, -NRxRx, -NHNRxRx, -NHC₁-C₃-alkyl-C(=O)Orx -NRxRc, where Rc is H, C₁-C₄ alkyl, -NRxRx; -C(=0)Rd, -CN, S(=O)pRx where Rd is Rd is C1-C4-alkyl, -ORx, -NRxRx C₁-C₃-alkyl-O-Cl-C3alkylC(=O)ORx, -C₁-C₃-alkyl-COORx; -C₁-C₃-alkyl-OH or C₁-C₄ alkyl ethers or esters thereof (O-C₁-C₃alkyl)q-O-Rx a 5 or 6 membered aromatic ring having 1-3 hetero atoms p is 1 or 2; Rx is independently selected from H, C₁-C₄ alkyl or acetyl; or a pair of Rx can together with the adjacent N atom form a ring; L is -0-, -S(=O), - or -CH₂-, where r is 0, 1 or 2; R₃-R₇ are substituents as defined in the specification; X is -(CR₈R₈)n-D-(CR₈R₈)m-; D is a bond, -NR₉-, -0-, -S-, -S(=O)- or -S(=O)₂-; and pharmaceutically acceptable salts and prodrugs thereof, have utility as HIV antivirals.

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